

Debate

Reinstatement of "germinal epithelium" of the ovary

Takashi Nishida*¹ and Naoyo Nishida²

Address: ¹Oita-Ken Saiseikai Hita Hospital, Social Welfare Organization, Saiseikai Imperial Gift Foundation Inc., 643-7 Miwa, Hita-shi, 877-1292, Japan and ²Department of Pathology, Kurume University School of Medicine, 67 Asahi-machi, Kurume-shi, 830-0011, Japan

Email: Takashi Nishida* - nojisan@saiseikai.hita.oita.jp; Naoyo Nishida - naoji@med.kurume-u.ac.jp

* Corresponding author

Published: 21 August 2006

Received: 10 August 2006

Reproductive Biology and Endocrinology 2006, **4**:42 doi:10.1186/1477-7827-4-42

Accepted: 21 August 2006

This article is available from: <http://www.rbej.com/content/4/1/42>

© 2006 Nishida and Nishida; licensee BioMed Central Ltd.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/2.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Background: The existing dogma that the former term ovarian "germinal epithelium" resulted from a mistaken belief that it could give rise to new germ cells is now strongly challenged.

Discussion: Two years ago, a research group of the University of Tennessee led by Antonin Bukovsky successfully demonstrated the oogenic process from the human ovarian covering epithelium now commonly called the ovarian surface epithelium. They showed the new oocyte with zona pellucida and granulosa cells, both originated from the surface epithelium arising from mesenchymal cells in the tunica albuginea, and stressed that the human ovary could form primary follicles throughout the reproductive period. This gives a big impact not only to the field of reproductive medicine, but also to the oncologic area. The surface epithelium is regarded as the major source of ovarian cancers, and most of the neoplasms exhibit the histology resembling müllerian epithelia. Since the differentiating capability of the surface epithelium has now expanded, the histologic range of the neoplasms in this category may extend to include both germ cell tumors and sex cord-stromal cell tumors.

Summary: Since the oogenic capability of ovarian surface cells has been proven, it is now believed that the oocytes can originate from them. The term "germinal epithelium", hence, might reasonably be reinstated.

Background

Unlike the avian ovary, where the follicles containing a large yolky egg look like a cluster of yellowish grapes, the human ovary is solid in structure, amygdaloid in shape, grayish white in color, and the follicles are usually invisible through the surface. Since the intraovarian follicle was firstly identified by Reinier De Graaf in 1672 [1], the origin of human oocyte has been disputed. Two opposing views have been known regarding the source of oocytes: (1) that they arise in the yolk sac [2-5], or (2) that they arise in the gonadal tissue itself [6-11]. The former view had gotten public consensus until the astonishing works

by Bukovsky et al. on the capability of ovarian covering tissue to produce new oocytes in adult human females were published [12-15].

A study of the histogenesis of ovarian components is essential to understand the oncogenesis of ovarian neoplasms. Since the ovarian covering tissue has now been revealed to have oogenic capability, the surface epithelium might be accountable as a source of germ cell tumors and sex cord-stromal cell tumors, as well as neoplasms exhibiting the müllerian histology.

Discussion

The nature of covering epithelium of the ovary is intriguing. Although only a part of peritoneal mesothelium, it proliferates to repair the minor trauma due to the ovulation, and thus has occasionally tumorigenic potential. The concept that the epithelium simulates the müllerian form in tumor formation has developed by degrees and now becomes a firm policy in the classification of ovarian cancer [16]. Although mucinous tumors are also placed in the common epithelial category, it is still disputed whether the origin of a certain group of mucinous neoplasms composed of intestinal type cells arose from germ cells, namely monophyletic teratomata.

In our previous experimental study, in which a chemical carcinogen, 7,12-dimethylbenz [a]anthracene (DMBA), was directly applied to the rat ovarian surface, an ovarian cancer was observed in about half of the DMBA-treated rats. The histology of the induced tumors also simulated the epithelia of rat genital tracts. In the experiment, however, an unexplainable cancer composed of heterologous osteoid tissue was observed [17]. At that time, if the covering tissue of the ovary were not called "surface epithelium", but named "germinal epithelium" instead, the tumor might have been classified as a teratomatous osteosarcoma.

Summary

From the capability of ovarian covering tissue to produce oocytes, the term "germinal epithelium" might be reinstated.

References

1. Houtzager HL: **Reinier De Graaf and his contribution to reproductive biology.** *Eur J Obstet Gynecol Reprod Biol* 2000, **90**:125-127.
2. Politzer G: **Über einen menschlichen Embryo mit 18 Ursegmentenpaar.** *Ztschr Anat Entwicklung* 1928, **87**:674-727.
3. Politzer G: **Über Zahl, Lage und Beschaffenheit der 'Urkeimzellen' eines menschlichen Embryo mit 26-27 Ursegmentpaaren.** *Ztschr Anat Entwicklung* 1928, **87**:766-80.
4. Politzer G: **Die Keimbahn des Menschen.** *Ztschr Anat Entwicklung* 1933, **100**:331-361.
5. Witschi E: **Migration of germ cell of human embryos from the yolk sac to the primitive gonadal folds.** *Contrib Embryol Carnegie Inst* 1948, **32**:67-80.
6. Waldeyer W: *Eierstock und Ei* Leipzig: Engelmann; 1870.
7. Allen E: **Ovogenesis during sexual maturity.** *Am J Anat* 1923, **31**:193-195.
8. Stieve H: **Die Entwicklung der Keimzellen und der Zwischenzellen in der Hodenanlage des Menschen.** *Ztschr mikr.-anat. Forsch* 1927, **10**:225-285.
9. Neumann HO: **Was wissen wir über die Keimbahn des Menschen?** *Arch Gynäk* 1929, **136**:107-144.
10. Evans HM, Swezy O: **Ovogenesis and the normal follicular cycle in adult mammalia.** *Mem Univ Calif* 1931, **9**:119-224.
11. Willis RA: **The borderland of embryology and pathology.** London: Butterworth; 1958.
12. Bukovsky A, Keenan JA, Caudle MR, Upadhyaya NB, Van Meter SE: **Immunohistochemical studies of the adult human ovary: possible contribution of immune and epithelial factors to folliculogenesis.** *Am J Reprod Immunol* 1995, **33**:323-340.
13. Bukovsky A, Caudle MR, Svetlikova M, Upadhyaya NB: **Origin of germ cells and formation of new primary follicles in adult human ovaries.** *Reprod Biol Endocrinol* 2004, **2**:20.

14. Bukovsky A, Caudle MR, Svetlikova M, Wimalasena J, Ayala ME, Dominguez R: **Oogenesis in adult mammals, including humans: a review.** *Endocrine* 2005, **26**:301-316.
15. Bukovsky A, Svetlikova M, Caudle MR: **Oogenesis in cultures derived from adult human ovaries.** *Reprod Biol Endocrinol* 2005, **3**:17.
16. Serov SF, Scully RE, Sobin LH: **Histological typing of ovarian tumors.** Geneva: World Health Organization; 1973.
17. Nishida T, Sugiyama T, Kataoka A, Ushijima K, Yakushiji M: **Histologic characterization of rat ovarian carcinoma induced by intraovarian insertion of a 7,12-dimethylbenz[a]anthracene-coated suture: common epithelial tumors of the ovary in rats?** *Cancer* 1998, **83**:965-970.

Publish with **BioMed Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours — you keep the copyright

Submit your manuscript here:
http://www.biomedcentral.com/info/publishing_adv.asp

